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# SPECIFICATION

Electronic Version 1.2.8

Stylesheet Version 1.0

## ELECTROLYTIC PROCESS AND APPARATUS

### Cross Reference to Related Applications

This application claims priority to U.S. Provisional Application Serial No. 60/345,307, filed on October 22, 2001, and U.S. Provisional Application Serial No. 60/369,090, filed on March 29, 2002, which are incorporated herein by reference in their entireties.

### Background of Invention

- [0001] This disclosure relates to electrolysis and more particularly, relates to improved electrolytic processes and apparatuses for the oxidation of inorganic or organic species.
- [0002] With the decline of gaseous chlorine as a microbiocide, various alternatives have been explored, including bleach, bleach with bromide, bromo-chlorodimethyl hydantoin, ozone, and chlorine dioxide ( $\text{ClO}_2$ ). Of these, chlorine dioxide has generated a great deal of interest for control of microbiological growth in a number of different industries, including the dairy industry, the beverage industry, the pulp and paper industries, the fruit and vegetable processing industries, various canning plants, the poultry industry, the beef processing industry and miscellaneous other food processing applications. Chlorine dioxide is also seeing increased use in municipal potable water treatment facilities and in industrial waste treatment facilities, because of its selectivity towards specific environmentally-objectionable waste materials, including phenols, sulfides, cyanides, thiosulfates, and mercaptans. In addition, chlorine dioxide is being used in the oil and gas industry for downhole applications as a well stimulation enhancement additive.
- [0003] Unlike chlorine, chlorine dioxide remains a gas when dissolved in aqueous

solutions and does not ionize to form weak acids. This property is at least partly responsible for the biocidal effectiveness of chlorine dioxide over a wide pH range, and makes it a logical choice for systems that operate at alkaline pH or that have poor pH control. Moreover, chlorine dioxide is a highly effective microbiocide at concentrations as low as 0.1 parts per million (ppm) over a wide pH range.

[0004] The biocidal activity of chlorine dioxide is believed to be due to its ability to penetrate bacterial cell walls and react with essential amino acids within the cell cytoplasm to disrupt cell metabolism. This mechanism is more efficient than other oxidizers that "burn" on contact and is highly effective against legionella, algae and amoebal cysts, giardia cysts, coliforms, salmonella, shigella, and cryptosporidium.

[0005] Unfortunately, chlorine dioxide in solution is unstable with an extremely short shelf life and thus, is not commercially available. Chlorine dioxide must typically be generated at its point of use such as, for example, by a reaction between a metal chlorate or metal chlorite in aqueous solution and a strong acid.

[0006] Electrochemical processes provide a means for generating chlorine dioxide for point of use applications. For example, U.S. Patent No. 5,419,816 to Sampson et al. describes a packed bed ion exchange electrolytic system and process for oxidizing species in dilute aqueous solutions by passing the species through an electrolytic reactor packed with a monobed of modified cation exchange material. A similar electrolytic process is described in U.S. Patent No. 5,609,742 to Sampson et al. for reducing species using a monobed of modified anion exchange.

[0007] One difficulty with electrochemical processes is that it can be difficult to control the generation of undesirable species. For example, there are many electrochemical reactions that can occur at the anode. Within a potential range of 0.90 to 2.10 volts, at least eight different reactions are thermodynamically possible, producing products such as chlorate ( $\text{ClO}_3^-$ ), perchlorate ( $\text{ClO}_4^-$ ), chlorous acid ( $\text{HClO}_2$ ), oxygen ( $\text{O}_2$ ), hydrogen peroxide ( $\text{H}_2\text{O}_2$ ) and ozone ( $\text{O}_3$ ). It is highly desirable and a significant commercial advantage to achieve high yield efficiency without producing these undesirable side reactions.

[0008] Chlorine dioxide has also been produced from a chlorine dioxide precursor

solution by contacting the precursor solution with a catalyst (e.g., catalysts containing a metal such as described for example in U.S. Pat. No. 5,008,096) in the absence of an electrical field or electrochemical cell. However, known catalytic processes have the disadvantage of becoming greatly deactivated within a matter of days. Moreover, it has been found that the support materials for the catalytic sites tend to quickly degrade due to the oxidizing nature of chlorine dioxide. Still further, the use of catalyst materials in packed columns or beds for generating chlorine dioxide has been found to cause a significant pressure drop across the column or form channels within the column that results in a significant decrease in conversion efficiency from the chlorine dioxide precursor to chlorine dioxide.

## Summary of Invention

- [0009] An electrolytic reactor for the electrocatalytic oxidation of chlorite ions in an aqueous solution comprises an anode; a cathode; and a particulate catalyst material.
- [0010] In another embodiment, an electrolytic reactor for the oxidation of chlorite ions in an aqueous solution comprises an anode compartment comprising an anode and a catalyst material; and a cathode compartment comprising a cathode.
- [0011] In another embodiment, an electrolytic reactor for the oxidation of chlorite ions in an aqueous solution comprises an anode compartment comprising an anode; a cathode compartment comprising a cathode; and a central compartment disposed between the anode and cathode compartments, wherein the central compartment comprises a catalyst material.
- [0012] In another embodiment, an electrolytic reactor for the oxidation of chlorite ions in an aqueous solution comprises an anode compartment comprising an anode; a cathode compartment comprising a cathode; and a central compartment disposed between the anode and cathode compartments, wherein the central compartment comprises a catalyst material, and wherein a non-permselective membrane separates the anode compartment from the central compartment.
- [0013] A process for fabricating a ceramic catalyst material comprises dissolving a metal oxide precursor into a solution; adding an additive consisting of anions or cations to the solution; coating a ceramic particle with the solution; and heating the coated

ceramic particle to a temperature effective to form a metal oxide.

- [0014] A process for generating chlorine dioxide from an alkali metal chlorite solution comprises applying a current to the electrolytic reactor, wherein the electrolytic reactor includes an anode, a cathode, and a catalyst material; flowing an aqueous alkali metal chlorite solution into the electrolytic reactor; and contacting the alkali metal chlorite solution with the catalyst material to electrocatalytically produce an effluent containing chlorine dioxide.
- [0015] In another embodiment, a process for generating chlorine dioxide from an alkali metal chlorite solution includes applying a current to an electrolytic reactor including an anode compartment comprising an anode and a catalyst material; and a cathode compartment comprising a cathode; flowing an aqueous alkali metal chlorite solution into the electrolytic reactor; and contacting the alkali metal chlorite solution with the catalyst material to electrocatalytically produce an effluent containing chlorine dioxide.
- [0016] In another embodiment, a process for generating chlorine dioxide from an alkali metal chlorite solution comprises applying a current to an electrolytic wherein the electrolytic reactor includes an anode compartment comprising an anode, a cathode compartment comprising a cathode, and a central compartment disposed between the anode and cathode compartments, wherein the central compartment comprises a catalyst material; flowing water into the anode compartment and generating hydrogen atoms; passing the hydrogen atoms into the central compartment; flowing the alkali metal chlorite solution into the central compartment; and contacting the alkali metal chlorite solution with the catalyst material to produce an effluent containing chlorine dioxide.
- [0017] In another embodiment, a process for generating chlorine dioxide from a dilute alkali metal chlorite solution comprises applying a current to an electrolytic reactor, wherein the electrolytic reactor comprises an anode compartment comprising an anode, a cathode compartment comprising a cathode, and a central compartment disposed between the anode and cathode compartments, wherein the central compartment comprises a catalyst material, and wherein a non-permselective membrane separates the anode compartment from the central compartment; flowing

water into the anode compartment and generating hydrogen atoms; passing the hydrogen atoms into the central compartment; flowing the dilute alkali metal chlorite solution into the central compartment; and contacting the dilute alkali metal chlorite solution with the catalyst material to produce an effluent containing chlorine dioxide.

[0018] The above described and other features will become better understood from the detailed description that is described in conjunction with the accompanying drawings.

### Brief Description of Drawings

[0019] Referring now to the figures wherein the like elements are numbered alike:

[0020] Figure 1 shows a cross sectional view illustrating a packed bed electrolytic reactor;

[0021] Figure 2 shows a partial cross sectional view illustrating the anode compartment of Figure 1, wherein the particulate material is layered;

[0022] Figure 3 shows a cross sectional view illustrating another packed bed electrolytic reactor;

[0023] Figure 4 shows a partial cross sectional view illustrating the central compartment of Figure 3, wherein the particulate material is layered;

[0024] Figure 5A and 5B show an exploded isometric view of an electrolytic reactor cassette;

[0025] Figure 6 is a graph showing a pressure drop of an electrolytic reactor as a function of time for different catalyst support materials;

[0026] Figure 7 is a graph showing conversion efficiency of an electrolytic reactor as a function of time during the electrolytic production of chlorine dioxide from an aqueous sodium chlorite feed solution;

[0027] Figure 8 is a graph showing a comparison of conversion efficiencies for different electrolytic reactors; and

[0028] Figure 9 is a graph showing a comparison of the applied voltage level as a function of time for different electrolytic reactors.



can be chosen to be non-permselective, e.g., a porous membrane. As used herein, the term "permselective" refers to a selective permeation of commonly charged ionic species through the membrane with respect to other diffusing or migrating ionic species having a different charge in a mixture. In contrast, the term "non-permselective" generally refers to a porous structure that does not discriminate among differently charged ionic species as the species pass through the porous structure, i.e., the membrane is non-selective with respect to ionic species. For example, in a permselective membrane such as a cation exchange membrane, cations can freely pass through the membrane whereas the passage of anions is prevented. In contrast, in a non-permselective membrane such as a porous membrane, the passage of anions and cations through the porous membrane are controlled by diffusion.

[0032] The cathode compartment 22 may or may not contain particulate material. If the cathode compartment 22 contains particulate material, as 24 shown in Figure 1, the cathode compartment 22 may be configured to possess its own physical properties (i.e., different properties from the particulate material 18 disposed in compartment 16) through which an aqueous solution can pass without entering adjacent compartment 16. Preferably, the particulate material 24 in the cathode compartment 22 comprises a cation exchange material.

[0033] An additive or additives, such as electrically conductive particles, may also be interspersed with the particulate materials 18, 24, to achieve certain results.

[0034] Referring now to Figure 3, there is shown a cross-sectional view illustrating an electrolytic reactor 100 including an anode compartment 102, a central compartment 104 and a cathode compartment 106. As shown, the central compartment 104 is interposed between the anode and cathode compartments 102, 106, respectively, and is separated from compartments 102, 106 by membranes 116, 118. As previously described, the membranes employed in the reactor may be a permselective membrane, a non-permselective membrane, or a combination of permselective and non-permselective membranes. In one embodiment, the membrane 116 separating the anode compartment 102 from the central compartment 104 is selected to be non-permselective. In yet another embodiment, the membranes 116, 118 separating the central compartment 104 from the adjacent anode and cathode compartments 102,







[0044] While the arrangements of anode, cathode, and packed bed electrolytic reactors illustrated in Figures 1, 2, and 3 are presently considered preferable; any arrangement in which a sufficient quantity of particulate material (including catalyst material) is packed between the anode and cathode in an electrolytic reactor or in at least one of the compartments of a divided electrolytic reactor can be used. Other embodiments include, but are not limited to, separation of the anode and cathode compartments to control intermixing of gases and solutions and provision of any number of packed-bed compartments separated by membranes placed between the anode and cathode to affect other oxidation, reduction or displacement reactions.

[0045] The anodes 12, 112 and the cathodes 14, 114 may be made of any suitable material based primarily on the intended use of the electrolytic reactor, costs and chemical stability. For example, the anode may be made of a conductive material, such as ruthenium, iridium, titanium, platinum, vanadium, tungsten, tantalum, oxides of at least one of the foregoing, combinations including at least one of the foregoing, and the like. Preferably, the anode comprises a metal oxide catalyst material disposed on a suitable support. The supports are typically in the form of a sheet, screen, or the like and are formed from a rigid material such as titanium, niobium, and the like. The cathode may be made from stainless steel, steel or may be made from the same material as the anode.

[0046] Permselective membranes, e.g., 20, 116, 118, preferably contain acidic groups so that ions with a positive charge can be attracted and selectively passed through the membrane in preference to anions. Preferably, the permselective membrane contains strongly acidic groups, such as  $\text{R-SO}_3^-$  and is resistant to oxidation and temperature effects. In a preferred embodiment, the permselective membranes are fluoropolymers that are substantially chemically inert to chlorine dioxide and the materials or environment used to produce the chlorine dioxide. Examples of suitable permselective membranes include perfluorosulfonate cation exchange membranes commercially available under the trade name NAFION commercially available from E.I. duPont de Nemours, Wilmington, DE.

[0047] Non-permselective membranes e.g., 20, 116, 118, contain pores that permit the non-selective diffusion of ionic species as well as non-ionic species from one



ions or the like that may then pass into the cathode compartment, if present. The use of the cation exchange material is especially useful when feeding a dilute alkali metal chlorite solution into compartment 16 or 104, as it helps lower the voltage within the compartment and increases conversion efficiency. The term "dilute" refers to aqueous alkali metal chlorite solutions containing less than about 10,000 milligrams alkali metal chlorite per liter of solution (mg/L), preferably less than about mg/L, and more preferably less than about 1,500 mg/L.

[0050] Examples of suitable cation exchange materials include, but are not intended to be limited to, polystyrene divinylbenzene cross-linked cation exchangers (e.g., strong acid types, weak acid types, iminodiacetic acid types, chelating selective cation exchangers and the like); strong acid perfluorosulfonated cation exchangers; naturally occurring cation exchangers, such as manganese greensand; high surface area macro-reticular or microporous type ion exchange resins having sufficient ion conductivity, and the like. For example, strong acid type exchange materials suitable for use are commercially available from Mitsubishi Chemical under the trade names Diaion SK116 and Diaion SK104. Optionally, the cation exchange material may be further modified, wherein a portion of the ionic sites are converted to semiconductor junctions, such as described in U.S. Patent Nos. 6,024,850, 5,419,816, 5,705,050 and 5,609,742, herein incorporated by reference in their entireties. In a preferred embodiment, the cation exchange materials have a cross-linking density greater than about 8 %, with greater than about 12 % more preferred and with greater than about 16 % even more preferred. Increasing the cross-linking density of the cation exchange materials has been found to increase the resistance of the cation exchange materials to oxidation and degradation. As a result, operating lifetimes for the electrolytic reactor can advantageously be extended during use for the production of strongly oxidizing products such as chlorine dioxide.

[0051] The particulate cation exchange material may also include electrically conductive particles, such as carbon and the like. The electrically conductive particles can be used to affect the transfer of DC current from the electrode to the membrane. For example, the use of electrically conductive particles can be used to lower the electrical resistance within the compartment. However, some additives, such as carbon, are prone to disintegration in acidic environments, thus requiring careful selection.

[0052] The packing density and conductivity of the cation exchange material within the compartment can be adjusted depending on the operating parameters and desired performance for the electrolytic reactors 10, 100, 130. For example, the cation exchange material may be shrunk before use in the electrolytic reactor, such as by dehydration or electrolyte adsorption. Dehydration may be by any method in which moisture is removed from the ion exchange material, for example, using a drying oven. It has been found that dehydration prior to packing can increase the packing density by as much as 40 %. Electrolyte adsorption involves soaking the material in a salt solution, such as sodium chloride. The packing density of the material so treated can be increased by as much as 20 %. The increase in packing density advantageously increases the volume in which the DC current travels, thus reducing the electrical resistance in the electrolytic reactor.

[0053] The catalyst material preferably comprises a support and an active metal catalyst. Preferably, the active metal catalyst is finely and discretely deposited onto the support. While not wanting to be bound by theory, it is believed that the active metal catalyst adsorbs hydrogen atoms without reacting with the hydrogen atoms, which form stable hydrides. Preferably, the active metal catalyst does not adsorb hydrogen atoms into its crystal lattice since this will decrease its catalytic activity. In a preferred embodiment, the active metal catalyst is a noble metal and does not react or dissolve with any of the components or solutions used in the electrolytic reactor 10, 100. While not wanting to be bound by theory, it is believed that the catalytic activity of the active metal is associated with crystal imperfections and the finely divided deposits help to increase the surface area as well as increase the number of active catalytic sites. Suitable active metal catalysts include, but are not limited to, ruthenium, platinum, palladium, osmium, iridium, rhodium, titanium, manganese, lead, zirconium, niobium, tantalum, tungsten, tin, and composites or mixtures or alloys or oxides of at least one of the foregoing metal catalysts. Preferably, the active metal catalyst is an oxide of a metal selected from the group consisting of ruthenium, platinum, palladium, osmium, iridium, and rhodium. More preferably, the active metal catalyst is a platinum oxide.

[0054] In another embodiment, the active metal catalyst is a combination of an oxide of ruthenium, platinum, palladium, osmium, iridium, rhodium or mixtures or alloys of at least one of the foregoing and a less active oxide of a metal including titanium, lead,

manganese, zirconium, niobium, tantalum, tungsten, tin or mixtures or alloys of at least one of the foregoing. Preferably, the molar ratio of the active metal catalyst to the less active metal catalyst is of about 0.3 : 1 to about 100 : 1. More preferably, the molar ratio of the active metal catalyst to the less active metal catalyst is about 10 : 1.

[0055] Preparation of the catalyst material generally includes contacting the support with a catalyst precursor to form the active metal catalyst sites, for example, a catalyst precursor salt. In one embodiment, a metal oxide precursor salt is dissolved in an aqueous solution including an alcohol, and the solution is coated onto the support. Suitable alcohols include methanol, ethanol, isopropanol, propanol, butanol, combinations including at least one of the foregoing alcohols, and the like. In a preferred embodiment, the metal oxide precursor salt is dissolved in isopropanol and deionized water solution. The amount of alcohol used in the solution is dependent on the solubility of the metal oxide precursor salt, wherein the volume fraction of alcohol in an aqueous solution is about 30 to about 90 % (v/v), with about 40 to about 80 % (v/v) preferred and with about 50 to about 60 % (v/v) more preferred.

[0056] Depending on the desired properties, a solution of the metal oxide precursor salt may contain further additives, for example, ions that increase the solubility of the metal oxide precursor. Suitable ions include hydroxides, chlorides, phosphates, sulfates, ammonium, potassium, sodium, lithium or the like. Preferably, the additive is ammonium hydroxide. The weight fraction of additive in the solution is about 0.1 to about 10 % (w/v), with about 0.5 to about 5 % (w/v) preferred and with about 1 % (w/v) more preferred.

[0057] Alternatively, the metal catalyst may be deposited onto the support material using other techniques such as impregnation, co-precipitation, ion exchange, dipping, spray coating, vacuum deposition, sputtering or the like. Preferably, the amount of metal catalyst deposited onto the support is about 0.01 to about 100-weight %, with about 0.01 to about 10-weight % more preferred, and with about 0.05 to about 2-weight % most preferred. In addition, it has been found that the catalyst activity of the catalyst material is improved with multiple depositions, i.e., the metal oxide precursor solution is applied by several individual depositions. Preferably, the number of depositions is about 2 to about 12, with the number of depositions of about 2 to about 3 more

preferred.

[0058] The metal deposited onto the support by the precursor solution is then thermally or chemically oxidized to the oxide form. When utilizing a thermal process, adjusting the temperature can control the extent of oxidation. The support is preferably thermally stable to the oxidation temperatures employed. The oxidation temperatures are preferably about 200 ° C to about 1000 ° C, with about 400 ° C to about 800 ° C more preferred, and with about 500 ° C to about 700 ° C even more preferred. Suitable chemical oxidants include permanganate. The extent of oxidation may be increased when a multiple deposition process is used in contrast to a single deposition process.

[0059] Suitable supports for the catalyst material include metals, zeolites, anthracite, glauconite, faujasite, mordenite, clinoptilolite, aluminas, silicas, clays, ceramics, carbon and the like. Of these supports, ceramics are most preferred. In a preferred embodiment, the catalyst materials are made from those ceramics described in U.S. Patents 4,725,390 and 4,632,876, herein incorporated by reference in their entireties. Preferred ceramics are those made essentially from nonmetallic minerals (such as mineral clays) by firing at an elevated temperature. More preferred are ceramic materials commercially available under the trade name MACROLITE<sup>®</sup> by the Kinetico Company. The MACROLITE<sup>®</sup> ceramic materials are spherically shaped and characterized by having a rough texture, high surface area, and level of moisture absorption of less than about 0.5%. The low level of moisture absorption allows for the metal oxide precursor solution to penetrate a minimal depth into the surface of the ceramic, thereby depositing metal onto the external surface of the support, an optimum location for subsequent contact with an aqueous solution. The surface area of the MACROLITE<sup>®</sup> ceramic materials is believed to be on the order of about 103 m<sup>2</sup> per gram.

[0060]

The process for making ceramic support materials, such as the MACROLITE<sup>®</sup> material, generally includes the following steps. In the first step, mineral particulate, binder, silicon carbide, and parting agent are mixed and spheroidized in order to form unfired spheroids. One example of mineral particulates contains: 60% orthoclase, 10% nepheline, 10% hornblende, 5% diopside, 15% accessory minerals (titanite, apatite,



magnetite and biotite) and trace amounts of secondary minerals (e.g. kaolinite and analcite). Another example contains approximately 75% plagioclase and orthoclase feldspar and 25% of the minerals pyroxene, hornblende, magnetite and quartz of which magnetite is less than 5%. Byproduct mineral fines of perlite (containing 2–5% chemically bound water) will also function as the mineral particulates. Minerals containing chemically bound water or sulfur, which are useful components of the mineral particulates, are hornblende, apatite, biotite, pyrite, vermiculite and perlite.

[0061] Binders that may be useful as raw materials include bentonite starch, polyvinyl alcohol, cellulose gum, polyvinyl acetate and sodium lignosulphonate. The amount of binder may generally comprise about 1 to about 5% by weight of the dry materials fed to the mixer and is generally sufficient to permit screening and handling of the spheroids without significant attrition or breakage.

[0062] Suitable parting agents include magnesium oxide, zircon, diaspore and high alumina clays as well as other surface metal oxides.

[0063] The spheroids are then dried at a temperature of about 40 ° C to about 200 ° C and typically screened. The ceramic spheroids are then over-fired in a kiln, which allows for the formation of the internal air cells, making the finished product less dense. The firing atmosphere is typically air. The silicon carbide in the spheroids is oxidized during firing, the SiC near the surface being more extensively oxidized than that in the core.

[0064] The product from the kiln is screened using standard methods known to those skilled in the art. Either before, during or after the screening step, the fired spheroids may be subjected to vigorous agitation by air or some other agitation means or to a water washing step in order to remove dust from their surfaces. The support material is then coated with an active metal catalyst in the manner previously described.

[0065] The particulate material, i.e., the cation exchange material or the catalyst material, is not intended to be limited to any particular shape. Suitable shapes include rods, extrudates, tablets, pills, irregular shaped particles, spheres, spheroids, capsules, discs, pellets or the like. In a preferred embodiment, the particulate material is spherical. More preferably, the particulate material includes a reticulated and textured

surface having an increased surface area. The size of the particulate material is dependent on the acceptable pressure drop across the bed of particulate material. The smaller the particulate material, the greater the pressure drop.

[0066] The size of the particulate material is generally defined by a mesh size of a sieve in which the particles are screened. For example, a 30 mesh sieve will allow particles less than about 600 micrometers to pass through the sieve, whereas particles about 600 micrometers or larger than will not pass through. Generally, sieves of varying mesh ratings are stacked and particles are separated using methods well known to those skilled in the art. The range of particles separated is defined by the mesh size of the sieves used. For example, a mesh size designation of 20/40 means that substantially all particles have a size ranging from 20 to 40 mesh (about 841 to about 420 micrometers in diameter). A 30/50 designation indicates that substantially all particles have a size ranging from 30 to 50 mesh (about 595 to about 297 micrometers in diameter). The sizes of the particulate matter are preferably smaller than 7 mesh (about 2800 micrometers in diameter), with smaller than 14 mesh (about 1400 micrometers in diameter) more preferred, and with smaller than 30 mesh (about 595 micrometers in diameter) even more preferred. Also preferred are particulate sizes greater than 60 mesh (about 250 micrometers in diameter), with greater than 50 mesh (about 297 micrometers in diameter) more preferred, and greater than 40 mesh (about 425 micrometers) even more preferred.

[0067] In a preferred mode of operation, a dilute aqueous feed solution of an alkali metal chlorite solution is passed through a selected compartment of the electrolytic reactor to generate an effluent containing chlorine dioxide. For example, in electrolytic reactor 100, the alkali metal chlorite solution preferably flows through central compartment 104. Water flows through the cathode and anode compartments 102, 106, respectively. Preferably, the water is deionized. As a current is applied to the reactor 100, the anode compartment 102 oxidizes the water to generate, among others, hydrogen ions whereas the cathode compartment 106 reduces the water to generate, among others, hydroxyl ions. The hydrogen ions generated in the anode compartment 102 can diffuse through membrane 116 into the central compartment 104. The hydrogen ions can be used to regenerate the cation exchange resin, if present, within the central compartment 104. Oxidation of the dissociated hydrogen

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[0068]

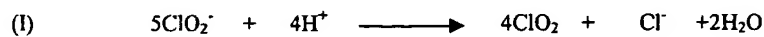
**[0069]**

[0070]

*(The following information was obtained from the FBI files maintained at the New York City Office.)*

[0071] In addition to temperature and current density, the contact time of the chlorine dioxide precursor with the catalyst material is preferably less than about 20 minutes and more preferably, less than about 2 minutes. Also preferred is a contact time greater than about 1 minute, with greater than about 0.1 minute more preferred. The velocity of the chlorine dioxide precursor solution through the catalyst bed is preferably less than about 100 centimeters/minute (cm/min), with less than about 70 cm/min more preferred and less than about 30 cm/min more preferred. Also preferred is a velocity greater than about 0.1 cm/min, with greater than about 10 cm/min more preferred and with greater than about 20 cm/min even more preferred. The pressure drop through the catalyst bed is preferably less than about 20 pounds per square inch (psi) and for most applications, with less than about 10 psi more preferred. Also preferred is a pressure drop greater than about 0.1 psi, and for most applications, with greater than about 1 psi more preferred. Further optimization for any of these process variables is well within the skill of those in the art in view of this disclosure.

[0072] While not wanting to be bound by theory, the use of the catalyst material in the electrochemical reactor system effectively lowers the activation energy for the oxidation of dissociated hydrogen ions into hydrogen atoms. Oxidation of the dissociated hydrogen ions into hydrogen atoms transpires so that the electrochemical oxidation of chlorite ions to chlorine dioxide can occur. It is believed that the electrochemical production of chlorine dioxide from chlorite ions is believed to proceed in accordance with the following reaction scheme:



[0073] Based on this reaction scheme, the maximum yield efficiency of chlorine dioxide will be an 80% conversion. Moreover, in addition to generating chlorine dioxide, the reaction advantageously produces as a byproduct, chloride ions. Chloride ions are not toxic or hazardous for most applications requiring the in situ generation of chlorine

dioxide.

[0074] In electrolytic reactors employing a non-permselective membrane to separate the anode compartment from the central compartment, the use of dilute chlorite solutions as described herein and the hygroscopic nature of the non-permselective membrane minimizes diffusion of the chlorite ions from the central compartment to the electrode compartments. Moreover, it is believed that employing cation exchange resins in the anode and/or cathode compartments further minimizes diffusion of chlorite ions from the central compartment to the respective electrode compartment. As a result, conversion efficiency of chlorite ion to chlorine dioxide is close to or about the maximum conversion efficiency.

### Example 1.

[0075] In this example, a ceramic catalyst material was prepared as follows. A metal oxide precursor solution was prepared by admixing 0.85 grams tetraamineplatinum (II) chloride, 41 ml of 91% isopropyl alcohol, 0.83 ml of 30% ammonium hydroxide and 26 ml of deionized water. The precursor solution was used twice to coat 100 milliliters of MACROLITE ML-20/40 that had been backwashed and air dried for about 12 hours. After each coating, the resin was baked at 550 °C for 30 minutes.

### Example 2.

[0076] In this example, a ceramic catalyst material was prepared as follows. A metal oxide precursor solution was prepared by admixing 0.85 grams tetraamineplatinum (II) chloride, 41 ml of 91% isopropyl alcohol, 0.83 ml of 30% ammonium hydroxide and 26 ml of deionized water. The precursor solution was used twice to coat 100 milliliters of SIR-600 resin commercially available from ResinTech, Inc. that had been backwashed and air-dried for about 12 hours. After each coating, the resin was baked at 550 °C for 30 minutes.

### Example 3.

[0077] In this example, electrochemical reactor cassettes were configured as described in Figures 4 and 5. The electrode compartments contained SK116 cation exchange resin commercially available from Mitsubishi Chemical. The central compartment contained



*[Faint, illegible text from bleed-through]*

$$\% \text{Yield} = \frac{\text{actual}}{\text{theoretical}} \times 100$$

wherein the actual yield is determined from the amount of chlorine dioxide generated, and wherein the theoretical yield is calculated by the amount of chlorine dioxide that could be generated from the sodium chlorite solution. As previously discussed, since it takes five moles of chlorite ions to make 4 moles of chlorine dioxide, the theoretical yield can be calculated as follows:

$$\%TheoreticalYield = \frac{[ClO_2]_{product}}{\left[ \frac{4}{5} [NaClO_2]_{feed} \right] \left[ \frac{90.5}{67.5} \right]} \times 100$$

wherein the term (90.5/67.5) is the ratio of the equivalent weight of the sodium chlorite to chlorine dioxide.

[0081] As shown in Figure 7, the conversion efficiency for the cassette employing the ceramic material is constant at about greater than about 95% of the maximum theoretical yield during the 1000 hours of continuous operation. Thus, the use of the catalyst material in the electrolytic reactor provides a yield that is at about the maximum possible yield. Increased conversion efficiencies over a prolonged period of time are a significant commercial advantage since it reduces the maintenance and operating costs of these reactors significantly.

### Example 4.

[0082]

In this example, two electrochemical reactor cassettes were configured as described in Figures 4 and 5. Each cassette included electrode compartments that contained SK116 cation exchange resin, wherein each compartment was separated from an adjacent compartment by cation exchange membranes (permselective). The central compartment of the first cassette contained a particulate material bed of three

equal layers. The first layer consisted of SK116 cation exchange resin; the second layer consisted of equal amounts by weight of the SK116 cation exchange resin and a ceramic catalyst material; and the third layer consisted of the ceramic catalyst material. The ceramic catalyst material was prepared in accordance with Example 1. The central compartment of the second cassette contained a packed bed of Diaion SK1B cation exchange resin. Commercially available from Mitsubishi Chemical, Diaion SK1B is a cross-linked polystyrene cation exchange resin having a cross-linking density of 8%.

[0083] Softened water was passed upwardly through the anode compartment at a flow rate of about 150 mL/min. A 25-weight % of sodium chlorite solution was added to the effluent of the anode compartment such that the final concentration of sodium chlorite was about 1000 mg/L. The combined sodium chlorite/effluent solution was then passed through the catalyst compartment. In the compartment including the three layers, the fluid passed from the bottommost layer to the topmost layer. In addition, softened water was passed upwardly through the cathode compartment at a flow rate of about 50 mL/min. While passing the solutions through the compartments of the reactor, a controlled current of about 4.0 amps was applied to the anode and cathode.

[0084] The results are summarized in Table II. All data recorded was taken after operating the electrolytic reactor for about 3 hours. The measured parameters for each cassette included cassette voltage, current, pressure drop through the bed flow rate (mL/min),  $\text{ClO}_2$  concentration (ppm), temperature (°C) and pH.

[t1]

Table II

	Catalyst Material	Cation Exchange Resin
$\text{ClO}_2$ concentration, (ppm)	558	11
Flow Rate, (mL/min)	150	158
Temperature, (°C)	26.5	26.5
pH	2.76	2.45
Pressure Drop, (psi)	8.0	13.0
Voltage, (volts)	23.0	19.2
Current, (Amps)	4.03	4.12



[0085] Clearly, the use of the ceramic catalyst material resulted in significantly superior results compared to the use of cation exchange resin in the central compartment. After only 3 hours of operation, conversion of chlorite ions to chlorine dioxide concentration was 50 times greater with the cassette employing the catalyst bed compared to the cassette employing the cation exchange resin bed. Moreover, the pressure drop across the bed increased significantly with the cassette employing the cation exchange resin indicating compaction of the bed.

### Example 5.

[0086] In this example, electrochemical reactor cassettes were configured as described in Figures 4 and 5. The cathode compartment was separated from the central compartment with a cation exchange membrane. The central compartment contained a particulate material bed of three equal layers. The first layer consisted of SK116 cation exchange resin; the second layer consisted of equal amounts by weight of the SK116 cation exchange resin and a catalyst material; and the third layer consisted of the catalyst material. The catalyst material was prepared in accordance with Example 2, i.e., modified SIR-600 resin. The electrode compartments contained SK106 cation exchange resin.

[0087] In one set of cassettes, the anode compartment was separated from the central compartment using a Flourtex<sup>®</sup> 9-70/22 non-permselective membrane commercially available from Sefar America, Inc. The Fluortex<sup>®</sup> is a fluorocarbon based membrane, and in this example, had a pore size of 0.070 millimeters. In a second set of cassettes, the anode compartment was separated from the central compartment using a cation exchange membrane.

[0088] The electrochemical reactor cassettes were operated in the following manner. Softened water having a conductivity of about 120  $\mu$  S/cm was passed upwardly through the anode compartment at a flow rate of about 150 mL/min. A 25-weight % sodium chlorite solution was added to the effluent of the anode compartment such that the final concentration of sodium chlorite was about 1000 mg/L. The combined sodium chlorite/effluent solution was then passed through the central compartment at a temperature of about 25 °C. In addition, softened water was passed upwardly through the cathode compartment at a flow rate of about 50 mL/min. While passing

the solutions through the various compartments of the reactor, a controlled current of about 4.0 amps was applied to the anode and cathode.

[0089] Figure 8 graphically illustrates the percent conversion of the chlorite ions to chlorine dioxide as a function of time for each set of cassettes. After an initial break-in period of about 200 hours of operation, the conversion efficiency for the cassettes that included the non-permselective membrane is greater than about 80-90% during the period of time monitored. Thus, the use of the non-permselective membrane between the anode compartment and the central compartment in the electrolytic reactor provided a yield that is at about the maximum possible yield. In contrast, the use of cassettes that included a cation exchange membrane (permselective) between the anode compartment and the central compartment exhibited a gradual decrease in conversion efficiency from about 1000 hours to about 2000 hours. Visual examination of the cation exchange membrane after about 2000 hours of operation indicated that oxidation had occurred. The cation exchange membrane exhibited discoloration, cracking, and delamination. In contrast, examination of the non-permselective membrane showed no visual evidence of oxidation.

[0090] Comparing the conversion efficiencies obtained in Figure 8 with the conversion efficiencies presented in Figure 7 illustrates the increased efficiency resulting from the use of the catalyst material prepared in accordance with Example 1 (ceramic media) as compared to the catalyst material in accordance with Example 2 (ion exchange resin).

[0091] Figure 9 graphically illustrates the applied voltage to the cassette as a function of time. The voltage was adjusted during operation of the electrolytic reactor to maintain a controlled current of about 4.0 amps. In the set of cassettes employing the non-permselective membrane, voltage was relatively constant throughout the 2000 hours of operation. In contrast, the voltage was gradually increased in the cassettes employing the cation exchange membrane (permselective). Since heterogeneous membranes such as the cation exchange membrane consist of colloidal ion exchange particles embedded in an inert binder, the gradual increase in voltage is believed to be caused by the oxidation of the ionthe ion exchange particles, which causes the material to swell within the rigid matrix, thereby decreasing the size of the interstices within the membrane and corresponding access to the ionic sites contained therein.

[0092] While the disclosure has been described with reference to an exemplary embodiment, it will be understood by those skilled in the art that various changes may be made and equivalents may be substituted for elements thereof without departing from the scope of the invention. In addition, many modifications may be made to adapt a particular situation or material to the teachings of the disclosure without departing from the essential scope thereof. Therefore, it is intended that the disclosure not be limited to the particular embodiment disclosed as the best mode contemplated for carrying out this disclosure, but that the disclosure will include all embodiments falling within the scope of the appended claims.